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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/807,512 04/08/2002		Peter I. Schrier	0652.2200000/EKS/SEZ	9121	
5	7590 02/11/2003				
Sterne Kessler Goldstein & Fox Suite 600 1100 New York Avenue NW			EXAMINER		
			DAVIS, MINH TAM B		
Washington, D	OC 20005-3934		ART UNIT	PAPER NUMBER	
			1642	1/	
			DATE MAILED: 02/11/2003		

Please find below and/or attached an Office communication concerning this application or proceeding.

	,	Application N	Application No.		Applicant(s)			
<del>(</del> ه	. Office Action Commence	09/807,512		SCHRIER ET AL.				
	Office Action Summary	Examiner		Art Unit				
		MINH-TAM DA		1642				
Period fo	The MAILING DATE of this communication app r Reply	ears on the cov	er sheet with the d	correspondence add	dress			
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).  - Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).  Status								
1) Responsive to communication(s) filed on 09 December 2002.								
2a) <u></u> ☐	This action is <b>FINAL</b> . 2b)⊠ Thi	is action is non	-final.					
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is								
closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213. <b>Disposition of Claims</b>								
4)⊠ Claim(s) <u>15-39</u> is/are pending in the application.								
4a) Of the above claim(s) is/are withdrawn from consideration.								
5) Claim(s) is/are allowed.								
6)□	6) Claim(s) is/are rejected.							
7)	Claim(s) is/are objected to.							
=	Claim(s) <u>15-39</u> are subject to restriction and/or	election requir	ement.					
	on Papers							
9) The specification is objected to by the Examiner.  10) The drawing(s) filed on is/are: a) □ accepted or b) □ objected to by the Examiner.								
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Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.								
If approved, corrected drawings are required in reply to this Office action.								
12) The oath or declaration is objected to by the Examiner.								
Priority under 35 U.S.C. §§ 119 and 120								
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).								
a) All b) Some * c) None of:								
1. Certified copies of the priority documents have been received.								
2. Certified copies of the priority documents have been received in Application No								
<ul> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>								
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).								
a) ☐ The translation of the foreign language provisional application has been received. 15)☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.								
Attachment	(s)							
2) Notice	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	4) [ 5) [ 6) [	Notice of Informal I	/ (PTO-413) Paper No( Patent Application (PTC				

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## **DETAILED ACTION**

## Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following 29 inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group 1, claim(s) 15, 34, drawn to the polypeptide of SEQ ID NO:2, and a method for inducing a cytotoxic T lymphocytes response *in vivo*, comprising administering the SEQ ID NO:2.

Groups 2-6, claim(s) 16-23, drawn to a CTL epitope comprising the amino acid sequence of SEQ ID NO:11, 12, 24, 25 or 26. Each amino acid sequence constitutes a single invention.

Groups 7-12, claims 24-33, drawn to a polynucleotide encoding SEQ ID NO:2, 11, 12, 24, 25 or 26. Each polynucleotide sequence constitutes a single invention.

Groups 13-17, claim 35, drawn a method for inducing a cytotoxic T lymphocytes response *in vivo*, comprising administering the SEQ ID NO: 11, 12, 24, 25 or 26. A method using each amino acid sequence constitutes a single invention.

Group 18, claim 36, drawn to a method for *ex vivo* treating an individual, comprising incubating CTL precursor cells from the individual with antigen presenting cells, and SEQ ID NO:2 and readministering said effector CTLs to the individual.

Groups 19-23, claim 37, drawn to a method for *ex vivo* treating an individual, comprising incubating CTL precursor cells from the individual with antigen presenting cells, and SEQ ID NO: 11, 12, 24, 25 or 26 and readministering said effector CTLs to the individual.

Group 24, claim 38, drawn to a method for *ex vivo* treating an individual, comprising administering to the individual cells transfected with the polynucleotide sequence encoding SEQ ID NO:2.

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Groups 25-29, claim 37, drawn to a method for *ex vivo* treating an individual, comprising administering to the individual cells transfected with the polynucleotide sequence encoding SEQ ID NO: 11, 12, 24, 25 or 26. A method using cells transfected with each of the polynucleotide sequence constitutes a single invention.

The inventions listed as Groups 1-29 do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

A national stage application shall relate to one invention only or to a group of inventions so linked as to form a single general inventive concept. When claims to different categories are present in the application, the claims will be considered to have unity of invention if the claims are drawn only to one of the following combinations of categories: (1) A product and a process specially adapted for the manufacture of said product; or (2) A product and a process of use of said product; or (3) A product, a process specially adapted for the manufacture of the said product, and a use of the said product; or (4) A process and an apparatus or means specifically designed for carrying out the said process; or (5) A product, a process specially adapted for the manufacture of the said product, and an apparatus or means specifically designed for carrying out the said process. If multiple products, processes of manufacture or uses are claimed, the first invention of the category first mentioned in the claims of the application will be considered as the main invention in the claims, see PCT article 17(3) (a) and 1.476 (c), 37 C.F.R. 1.475(b) and (d). Group 1 will be the main invention. After that, all other products and methods will be broken out as separate groups (see 37 CFR 1.475(d).)

Group 1, claims 15, 34 form a single general inventive concept.

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Groups 2-12 are not linked to the single general inventive concept of Group 1, because they are additional products which do not share the same structure with the polypeptide of SEQ ID NO:2 of group 1.

Group 18 is an additional use claimed for the polypeptide of SEQ ID NO:2 of group 1.

Groups 13-17, 19-23 are not linked to the single general inventive concept of Group 1, because they are additional methods which do not recite the use of the polypeptide of SEQ ID NO:2 of group 1.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to MINH-TAM DAVIS whose telephone number is 703-305-2008. The examiner can normally be reached on 9:30AM-4:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, ANTHONY CAPUTA can be reached on 703-308-3995. The fax phone

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numbers for the organization where this application or proceeding is assigned are 703-872-9306 for regular communications and 703-872-9307 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0916.

MINH TAM DAVIS

February 7, 2003

ANTHONY C. COPUTA
CUPERVISORY PATENT EXAMINER
TECHNOLOGY CENTER (800)

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